



(11) **EP 0 961 174 B1**

(12) **EUROPEAN PATENT SPECIFICATION**

(45) Date of publication and mention  
of the grant of the patent:  
**03.01.2007 Bulletin 2007/01**

(51) Int Cl.:  
**G03F 7/20** <sup>(2006.01)</sup> **G03F 1/14** <sup>(2006.01)</sup>  
**B01J 19/00** <sup>(2006.01)</sup>

(21) Application number: **99110362.3**

(22) Date of filing: **28.05.1999**

(54) **Compositions and methods involving direct write optical lithography**

Zusammensetzungen und Verfahren für direkt schreibende optische Lithographie

Compositions et méthodes pour la lithographie optique à écriture directe

(84) Designated Contracting States:  
**AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU**  
**MC NL PT SE**  
Designated Extension States:  
**AL LT LV MK RO SI**

(30) Priority: **29.05.1998 US 87333 P**  
**26.05.1999 US 318775**

(43) Date of publication of application:  
**01.12.1999 Bulletin 1999/48**

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**WO-A-99/42813** **WO-A-99/63385**  
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- **MARK DAVIDSON : "A MICROLENS DIRECT-WRITE CONCEPT FOR LITHOGRAPHY"**  
**PROCEEDINGS OF THE SPIE**, vol. 3048, 1997,  
pages 346-355, XP000866011 US
- **R. SELTMANN ET AL.: "New system for fast submicron optical direct writing"**  
**MICROELECTRONIC ENGINEERING**, vol. 30, no. 1, 1996, pages 123-127, XP004003046 amsterdam, nl ISSN: 0167-9317
- **DAVID W. MONK & RICHARD O. GALE : "The Digital Micromirror Device for Projection Display"**  
**MICROELECTRONIC ENGINEERING**, vol. 27, no. 1, 1 February 1995 (1995-02-01), pages 489-493, XP004025127 amsterdam, NL ISSN: 0167-9317

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## Description

### BACKGROUND OF THE INVENTION

#### Technical Field of the Invention

**[0001]** This invention relates to optical lithography and more particularly to direct write optical lithography.

#### Description of the Related Art

**[0002]** Polymer arrays, such as the GeneChip® probe array (Affymetrix, Inc., Santa Clara, CA), can be synthesized using light-directed methods described, for example, in U.S. Patent Nos. 5,143,854; 5,424,186; 5,510,270; 5,800,992; 5,445,934; 5,744,305; 5,384,261 and 5,677,195 and PCT published application no. WO 95/11995. As an example, light-directed synthesis of oligonucleotides employs 5'-protected nucleosidephosphoramidite "building blocks." The 5'-protecting groups may be either photolabile or acid-labile. A plurality of polymer sequences in predefined regions are synthesized by repeated cycles of deprotection (selective removal of the protective group) and coupling. Coupling (i.e., nucleotide or monomer addition) occurs only at sites that have been deprotected. Three methods of light-directed synthesis are: use of photolabile protecting groups and direct photodeprotection (DPD); use of acid-labile 4,4'-dimethoxytrityl (DMT) protecting groups and a photoresist; use of DMT protecting groups and a polymer film that contains a photoacid generator (PAG).

**[0003]** These methods have many process steps similar to those used in semiconductor integrated circuit manufacturing. These methods also often involve the use of photomasks (masks) that have a predefined image pattern which permits the light used for synthesis of the polymer arrays to reach certain regions of a substrate but not others. The substrate can be non-porous, rigid, semi-rigid, etc. It can be formed into a well, a trench, a flat surface, etc. The substrate can include solids, such as siliceous substances such as silicon, glass, fused silica, quartz, and other solids such as plastics and polymers, such as polyacrylamide, polystyrene, polycarbonate, etc. Typically, the solid substrate is called a wafer from which individual chips are made. As such, the pattern formed on the mask is projected onto the wafer to define which portions of the wafer are to be deprotected and which regions remain protected. See, for example, U.S. Patent Nos. 5,593,839 and 5,571,639.

**[0004]** The lithographic or photochemical steps in the synthesis of nucleic acid arrays may be performed by contact printing or proximity printing using photomasks. For example, an emulsion or chrome-on-glass mask is placed in contact with the wafer, or nearly in contact with the wafer, and the wafer is illuminated through the mask by light having an appropriate wavelength. However, masks can be costly to make and use and are capable of being damaged or lost.

**[0005]** In many cases a different mask having a particular predetermined image pattern is used for each separate photomasking step, and synthesis of a wafer containing many chips requires a plurality of photomasking steps with different image patterns. For example, synthesis of an array of 20mers typically requires approximately seventy photolithographic steps and related unique photomasks. So, using present photolithographic systems and methods, a plurality of different image pattern masks must be pre-generated and changed in the photolithographic system at each photomasking step. This plurality of different pattern masks adds lead time to the process and complexity and inefficiency to the photolithographic system and method. Further, contact printing using a mask can cause defects on the wafer so that some of the reaction sites are rendered defective. Thus, a photolithographic system and method that does not require such masks and obviates such difficulties would be generally useful in providing a more efficient and simplified lithographic process.

**[0006]** U.S. Patent No. 5 424 186 discloses a method of synthesizing nucleotides on a substrate by deprotecting reaction sites on a substrate, wherein light from a light source is directed to the substrate through a mask. Alternatively, a method is disclosed using liquid crystals (i.e. "light valves") to selectively deprotect reaction sites on the substrate.

**[0007]** The article SPIE, Vol. 3048, pp. 346-355 (1997) (XP866011) by Mark Davidson, discloses use of arrays of microlenses for direct-write photolithography on wafers. However, this article does not disclose to provide a predetermined two dimensional light pattern four deprotecting protected reaction sites on a substrate.

**[0008]** The article Microelectronic Engineering, Vol. 30, pp. 123-127 (1996) (XP4003046) by R. Selmann et al. discloses a method for direct-write photolithography on wafers, utilizing phase modulating spatial light modulator for imaging laser light onto the wafer. The spatial light modulator employs a a CMOS active matrix and a reflective, viscoelastic layer on top. However, this article does not disclose to provide a predetermined two dimensional light pattern for deprotecting protected reaction sites on substrate.

**[0009]** The article Microelectronic Engineering, Vol. 27, pp. 489-493 (1995) (XP4025127) by D. Monk et al. discloses a digital micro-mirror device utilizing an array of micro-mirrors, each micro-mirror being mounted on a hinge over an integrated circuit structure operable to vary the orientation of the micro-mirror. However, this article does not disclose to provide a predetermined two dimensional light pattern for deprotecting protected reaction sites on a substrate.

### SUMMARY OF THE INVENTION.

**[0010]** The invention provides methods for deprotecting protected reaction sites on a substrate as defined in claims 1-7, and optical lithography systems for depro-

tecting reaction sites on a substrate as defined in claims 8-11.

**[0011]** In view of the above, one advantage of the invention is providing an improved and simplified system and method for optical lithography for deprotecting protected reaction sites on a substrate.

**[0012]** Another advantage of the present invention is providing an optical lithography system and method that dynamically generates an image using a computer and reconfigurable spatial light modulator, employing an array of deformable membrane mirrors.

**[0013]** A further advantage of the present invention is providing an optical lithography system and method that does not use photomasks.

**[0014]** A still further advantage of the present invention is providing an optical lithography system and method that uses computer generated electronic control signals and a spatial light modulator, without any photomask, to project a predetermined light pattern onto a surface of a substrate for the purposes of deprotecting various areas of a polymer array.

**[0015]** In connection with the invention, polymer array synthesis is performed using a system without photomasks.

**[0016]** In connection with the invention, polymer array synthesis is performed using a system with a transmissive spatial light modulator and without a lens and photomask

**[0017]** In connection with the invention, a Direct Write System transmits image patterns to be formed on the surface of a substrate (e.g., a wafer). The image patterns are stored in a computer. The Direct Write System projects light patterns generated from the image patterns onto a surface of the substrate for light-directed polymer synthesis (e.g., oligonucleotide). The light patterns are generated by a spatial light modulator controlled by a computer, rather than being defined by a pattern on a photomask. Thus, in the Direct Write System each pixel is illuminated with an optical beam of suitable intensity and the imaging (printing) of an individual feature on a substrate is determined dynamically by computer control.

**[0018]** In connection with the invention, polymer array synthesis is accomplished using a class of devices known as spatial light modulators employing an array of deformable membrane mirrors to define the image pattern of the polymer array to be deprotected.

**[0019]** An even further aspect of the present invention provides methods for synthesizing polymer arrays using spatial light modulators and the polymer arrays synthesized using the methods taught herein.

**[0020]** As can be appreciated by one skilled in the art, the invention is relevant to optical lithography in general, and more specifically to optical lithography for polymer array synthesis using photolithographic processes. However, it is inherent that the invention is generally applicable to eliminating the need for a photomask in optical lithography.

## BRIEF DESCRIPTION OF THE DRAWINGS

**[0021]** The above objects, features, and advantages of the present invention will become more apparent from the following detailed description taken with the accompanying drawings in which:

Figure 1 shows a first embodiment of the invention having a light source, a reflective spatial light modulator, such as a micro-mirror array, and a lens.

Figure 2 is a diagrammatic representation of a second embodiment of the invention employing an array of, for example, micro-lenses.

Figure 3 illustrates a micro-lens array in the form of Fresnel Zone Plates, which may be used in the invention.

Figure 4 shows a third embodiment of the invention having a transmissive spatial light modulator.

## DESCRIPTION OF THE PREFERRED EMBODIMENTS

**[0022]** The presently preferred invention is based on the principle that a Direct Write Optical Lithography System will significantly improve the cost, quality, and efficiency of polymer array synthesis by providing a maskless optical lithography system and method where predetermined image patterns can be dynamically changed during photolithographic processing. As such, an optical lithography system is provided to include a means for dynamically changing an intended image pattern without using a photomask. One such means includes a spatial light modulator that is electronically controlled by a computer to generate unique predetermined image patterns at each photolithographic step in polymer array synthesis. The spatial light modulators can be, for example, micromachined mechanical modulators or microelectronic devices (e.g. liquid crystal display (LCD)). The Direct Write System of the present invention using such spatial light modulators is particularly useful in the synthesis of polymer arrays, such as polypeptide, carbohydrate, and nucleic acid arrays. Nucleic acid arrays typically include polynucleotides or oligonucleotides attached to glass, for example, Deoxyribonucleic Acid (DNA) arrays.

**[0023]** Certain preferred embodiments of the invention involve use of the micromachined mechanical modulators to direct the light to predetermined regions (i.e., known areas on a substrate predefined prior to photolithography processing) of the substrate on which the polymers are being synthesized. The predetermined regions of the substrate associated with, for example, one segment (referred to herein as a pixel) of a micromachined mechanical modulator (e.g., a micro-mirror array) are referred to herein as features. In each predetermined region or feature a particular oligonucleotide sequence, for example, is synthesized. The mechanical modulators come in a variety of types, two of which will be discussed in some detail below.

**[0024]** One type of mechanical modulator is a micro-

mirror array which uses small metal mirrors to selectively reflect a light beam to particular individual features; thus causing the individual features to selectively receive light from a light source (i.e., turning light on and off of the individual features). An example is the programmable micro-mirror array Digital Micromirror Device (DMD™) manufactured by Texas Instruments, Inc., Dallas, Texas, USA. Texas Instruments markets the arrays primarily for projection display applications (e.g., big-screen video) in which a highly magnified image of the array is projected onto a wall or screen. The present invention shows, however, that with appropriate optics and an appropriate light source, a programmable micro-mirror array can be used for photolithographic synthesis, and in particular for polymer array synthesis.

**[0025]** The Texas Instruments' DMD™ array consists of 640 x 480 mirrors (the VGA version) or 800 x 600 mirrors (the super VGA (SVGA) version). Devices with more mirrors are under development. Each mirror is 16 μm x 16 μm and there are μm gaps between mirrors. The array is designed to be illuminated 20 degrees off axis. Each mirror can be turned on (tilted 10 degrees in one direction) or off (tilted 10 degrees in the other direction). A lens (on axis) images the array onto a target. When a micro-mirror is turned on, light reflected by the micro-mirror passes through the lens and the image of the micro-mirror appears bright. When a micro-mirror is turned off, light reflected by the micro-mirror misses the lens and the image of the micro-mirror appears dark. The array can be reconfigured by software (i.e., every micro-mirror in the array can be turned on or off as desired) in a fraction of a second.

**[0026]** An optical lithography system including a micro-mirror array 1 based spatial light modulator according to one embodiment of the invention is shown in FIG. 1. This embodiment includes a spatial light modulator made of a micro-mirror array 1, and arc lamp 3, and a lens 2 to project a predetermined image pattern on a chip or wafer (containing many chips) 4. In operation, collimated, filtered and homogenized light 5 from the arc lamp 3 is selectively reflected as a light beam 6 according to dynamically turned on micro mirrors in the micro-mirror array 1 and transmitted through lens 2 on to chip or wafer 4 as reflected light beam 8. Reflected light from micro-mirrors that are turned off 7 is reflected in a direction away from the lens 2 so that these areas appear dark to the lens 2 and chip or wafer 4. Thus, the spatial light modulator, micro-mirror array 1, modulates the direction of reflected light (6 and 7) so as to define a predetermined light image 8 projected onto the chip or wafer 4. The direction of the reflected light alters the light intensity transmitted from each pixel to each feature. In essence, the spatial light modulator operates as a directional and intensity modulator.

**[0027]** The micro-mirror array 1 can be provided by, for example, the micro-mirror array of the Texas Instruments(TI) DMD, in particular, the TI "SVGA DLP™" subsystem. The Texas Instruments "SVGA DLP™" subsystem

with optics may be modified for use in the present invention. The Texas Instruments "SVGA DLP™" subsystem includes a micro-mirror array (shown as micro-mirror array 1 in Fig. 1), a light source, a color filter wheel, a projection lens, and electronics for driving the array and interfacing to a computer. The color filter wheel is replaced with a bandpass filter having, for example, a bandpass wavelength of 365-410 nm (wavelength dependent upon the type of photochemicals selected for used in the process). For additional brightness at wavelengths of, for example, 400-410 nm, the light source can be replaced with arc lamp 3 and appropriate homogenizing and collimating optics. The lens included with the device is intended for use at very large conjugate ratios and is replaced with lens 2 or set of lenses appropriate for imaging the micro-mirror array 1 onto chip or wafer 4 with the desired magnification. Selection of the appropriate lens and bandpass filter is dependent on, among other things, the requisite image size to be formed on the chip, the type of spatial light modulator, the type of light source, and the type of photoresist and photochemicals being used in the system and process.

**[0028]** A symmetric lens system (e.g., lenses arranged by type A-B-C-C-B-A) used at 1:1 magnification (object size is the same as the image size) is desirable because certain aberrations (distortion, lateral color, coma) are minimized by symmetry. Further, a symmetric lens system results in a relatively simple lens design because there are only half as many variables as in an asymmetric system having the same number of surfaces. However, at 1:1 magnification the likely maximum possible chip size is 10.88 mm x 8.16 mm with a VGA device, or 10.2 mm x 13.6 mm with an SVGA device. Synthesis of, for example, a standard GeneChip® 12.8 mm x 12.8 mm chip uses an asymmetric optical system (e.g., a magnification of about 1.25:1 with SVGA device) or a larger micro-mirror array (e.g., 1028 x 768 mirrors) if the mirror size is constant. In essence, the lens magnification can be greater than or less than 1 depending on the desired size of the chip.

**[0029]** In certain applications of the invention, a relatively simple lens system, such as a back-to-back pair of achromats or camera lens, is adequate. A particularly useful lens for some applications of the invention is the Rodenstock (Rockford, IL) Apo-Rodagon D. This lens is optimized for 1:1 imaging and gives good performance at magnifications up to about 1.3:1. Similar lenses may be available from other manufacturers. With such lenses, either the Airy disk diameter or the blur circle diameter will be rather large (maybe 10 μm or larger). See *Modern Optical Engineering*, 2d Edition, Smith, W.J., ed., McGraw-Hill, Inc., New York (1990). For higher-quality synthesis, the feature size is several times larger than the Airy disk or blur circle. Therefore, a custom-made lens with resolution of about 1-2 μm over a 12.8 mm x 12.8 mm field is particularly desirable.

**[0030]** A preferred embodiment of synthesizing polymer arrays with a programmable micro-mirror array using

the DMT process with photoresist takes place as follows. First, a computer file is generated and specifies, for each photolithography step, which mirrors in the micro-mirror array 1 need to be on and which need to be off to generate a particular predetermined image pattern. Next, the individual chip or the wafer from which it is made 4 is coated with photoresist on the synthesis surface and is mounted in a holder or flow cell (not shown) on the photolithography apparatus so that the synthesis surface is in the plane where the image of the micro-mirror array 1 will be formed. The photoresist may be either positive or negative thus allowing deprotection at locations exposed to the light or deprotection at locations not exposed to the light, respectively (example photoresists include: negative tone SU-8 epoxy resin (Shell Chemical) and those shown in the above cited patents and U.S. patent appl. no. 08/634,053, now U.S. Patent No. 5 959 098, issued September 28, 1999). A mechanism for aligning and focusing the chip or wafer is provided, such as a x-y translation stage. Then, the micro-mirror array 1 is programmed for the appropriate configuration according to the desired predetermined image pattern, a shutter in the arc lamp 3 is opened, the chip or wafer 4 is illuminated for the desired amount of time, and the shutter is closed. If a wafer (rather than a chip) is being synthesized; a stepping-motor-driven translation stage moves the wafer by a distance equal to the desired center-to-center distance between chips and the shutter of the arc lamp 3 is opened and closed again, these two steps being repeated until each chip of the wafer has been exposed.

**[0031]** Next, the photoresist is developed and etched. Exposure of the wafer 4 to acid then cleaves the DMT protecting groups from regions of the wafer where the photoresist has been removed. The remaining photoresist is then stripped. Then DMT-protected nucleotides containing the desired base (adenine (A), cytosine (C), guanine (G), or thymine (T)) are coupled to the deprotected oligonucleotides.

**[0032]** Subsequently, the chip or wafer 4 is re-coated with photoresist. The steps from mounting the photoresist coated chip or wafer 4 in a holder through re-coating the chip or wafer 4 with photoresist are repeated until the polymer array synthesis is complete.

**[0033]** It is worth noting that if a DPD method, using for example 1-(6-nitro-1,3-benzodioxol-5-yl)ethoxycarbonyl (MeNPOC) chemistry, or a PAG method, using a polymer film containing a photoacid generator (PAG), are used for polymer array synthesis then photoresist would not be used and the process is somewhat simplified. However, the use of a direct write optical lithography system with a spatial light modulator is also applicable to performing a process of deprotection of reaction sites using the DPD and PAG methods without photoresist.

**[0034]** As is clear from the above described method for polymer array synthesis, no photomasks are needed. This simplifies the process by eliminating processing time associated with changing masks in the optical lithography system and reduces the manufacturing cost for polymer

array synthesis by eliminating the cost of the masks as well as processing defects associated with using masks. Therefore, the optical lithography system is simplified and more efficient than conventional photomask based optical lithography systems. This is particularly valuable in complex multiple step photolithography processing; for example polymer array synthesis of GeneChip<sup>®</sup> probe arrays having upwards of seventy or more cycles, especially when many different products are made and revised.

**[0035]** As indicated above, substrates coated with photoresist are employed in preferred embodiments of the invention, e.g., using the DMT process with photoresist. The use of photoresist with photolithographic methods for fabricating polymer arrays is discussed in McGall et al., *Chemtech*, pp. 22-32 (February 1997); McGall et al., *Proc. Natl. Acad. Sci., U.S.A.*, Vol. 93, pp. 13555-13560 (Nov. 1996) and various patents cited above. Alternatively, polymer array synthesis processing can be performed using photoacid generators without using a conventional photoresist, e.g., using the PAG process, or using direct photodeprotection without using any photoresist, e.g., using the DPD process. The use of photoacid generators is taught in U.S. Application No. 08/969,227, filed November 13, 1997, now U.S. Patent No. 6 083 697, issued July 4, 2000. However, the present invention is particularly useful when using the DMT and PAG processes for polymer array synthesis.

**[0036]** When synthesizing nucleic acid arrays, the photochemical processes used to fabricate the arrays is preferably activated with light having a wavelength greater than 365 nm to avoid photochemical degradation of the polynucleotides used to create the polymer arrays. Other wavelengths may be desirable for other probes. Many photoacid generators (PAGs) based on *o*-nitrobenzyl chemistry are useful at 365 nm. Further, when using the mirror array from Texas Instruments discussed above, the PAG is preferably sensitive above 400 nm to avoid damage to the mirror array. To achieve this, *p*-nitrobenzyl esters can be used in conjunction with a photosensitizer. For example, *p*-nitrobenzyltosylate and 2-ethyl-9,10-dimethoxy-anthracene can be used to photochemically generate toluenesulfonic acid at 405 nm. See S.C. Busman and J.E. Trend, *J. Imag. Technol.*, 1985, 11, 191; A. Nishida, T. Hamada, and O. Yonemitsu, *J. Org. Chem.*, 1988, 53, 3386. In this system, the sensitizer absorbs the light and then transfers the energy to the *p*-nitrobenzyltosylate, causing dissociation and the subsequent release of toluenesulfonic acid. Alternate sensitizers, such as pyrene, *N,N*-dimethylnaphthylamine, perylene, phenothiazine, canthone, thiocanthone, actophenone, and benzophenone that absorb light at other wavelengths are also useful.

**[0037]** A variety of photoresists sensitive to 436-nm light are available for use in polymer array synthesis and will avoid photochemical degradation of the polynucleotides.

**[0038]** A second preferred mechanical modulator that

may be used in the invention is the Grating Light Valve™ (GLV™) available from Silicon LightMachines, Sunnyvale, CA, USA. The GLV™ relies on micromachined pixels that can be programmed to be either reflective or diffractive (Grating Light Valve™ technology). Information regarding certain of the mechanical modulators discussed herein is obtained at <http://www.ti.com> (Texas Instruments) and <http://siliconlight.com>. (Silicon LightMachines).

**[0039]** Although preferred spatial light modulators include the mechanical modulators DMD™ available from Texas Instruments and the GLV™ available from Silicon LightMachines, various types of spatial light modulators exist and may be used in the practice of the present invention. See *Electronic Engineers' Handbook*, 3<sup>rd</sup> Ed., Fink, D.G. and Christiansen, D. Eds., McGraw-Hill Book Co., New York (1989). Deformable membrane mirror arrays are available from Optron Systems Inc. (Bedford, MA). Liquid-crystal spatial light modulators are available from Hamamatsu (Bridgewater, NJ), Spatialight (Novato, CA), and other companies. However, one skilled in the art must be careful to select the proper light source and processing chemistries to ensure that the liquid-crystal spatial light modulator is not damaged since these devices may be susceptible to damage by various ultraviolet (UV) light. Liquid-crystal displays (LCD; e.g., in calculators and notebook computers) are also spatial light modulators useful for photolithography particularly to synthesize large features. However, reduction optics would be required to synthesize smaller features using LCDs.

**[0040]** Some spatial light modulators may be better suited than the Texas Instruments device for use with UV light and would therefore be compatible with a wider range of photoresist chemistries. One skilled in the art will choose the spatial modulator that is compatible with the chosen wavelength of illumination and synthesis chemistries employed. For example, the device from Texas Instruments DMD™ should not be used with UV illumination because its micro-mirror array may be damaged by UV light. However, if the passivation layer of the micro-mirror array is modified or removed, the Texas Instruments DMD™ could be used in the invention with UV light.

**[0041]** One embodiment that is particularly useful when extremely high resolution is required involves imaging the micro-mirror array using a system of the type shown in Fig. 2. In this system, a lens 12 images the micro-mirror array 11 (e.g., DMD™ or GLV™) onto an array 10 having an array of micro-lenses 15 or non-imaging light concentrators. Each element of the array 10 focuses light onto the chip or wafer, e.g., Gene Chip array 14. Each micro-lens 15 produces an image of one pixel of the micro-mirror array 11. Optics 16, including a shaping lens 17 may be included to translate light from a light source 13 onto the micro-mirror array 11.

**[0042]** For example, if an SVGA DLP™ device is imaged with 1:1 magnification onto a micro-lens array 10, an appropriate micro-lens array 10 can consist of 800 x

600 lenses (micro-lenses 15) with 17  $\mu\text{m}$  center-to-center spacing. Alternatively, the micro-lens array can consist of 400 x 300 17  $\mu\text{m}$  diameter lenses with 34  $\mu\text{m}$  center-to-center spacing, and with opaque material (e.g., chrome) between micro-lenses 15. One advantage of this alternative is that cross-talk between pixels is reduced. The light incident upon each micro-lens 15 can be focused to a spot size of 1-2  $\mu\text{m}$ . Because the spot size is much less than the spacing between micro-lenses, a 2-axis translation stage (having, in these examples, a range of travel of at least either 17  $\mu\text{m}$  x 17  $\mu\text{m}$  or 34  $\mu\text{m}$  x 34  $\mu\text{m}$ ) is necessary if complete coverage of the chip or wafer 14 is desired.

**[0043]** Micro-lenses 15 can be diffractive, refractive, or hybrid (diffractive and refractive). Refractive micro-lenses can be conventional or gradient-index. A portion of a diffractive micro-lens array 10 is shown in Figure 3 and has individual micro-lenses formed as circles commonly known as Fresnel Zone Plates 20. Alternatively an array of non-imaging light concentrators can be employed. An example of such an approach would include a short piece of optical fiber which may be tapered to a small tip.

**[0044]** Furthermore, some spatial light modulators are designed to modulate transmitted rather than reflected light. An example of a transmissive spatial light modulator is a liquid crystal display (LCD) and is illustrated in another embodiment, shown in Fig. 4. This embodiment includes a light source 33 providing light 35, transmissive spatial light modulator 31 and a computer 39 providing electronic control signals to the transmissive spatial light modulator 31 through cables 40 so as to transmit a desired light image 38 on the chip or wafer 34. The computer 39 may be, for example, a unique programmable controller, a personal computer (PC), or a CAD system used to design the desired image pattern.

**[0045]** Using a transmissive spatial light modulator has even additional advantages over the conventional optical lithography system. Reflective spatial light modulators require a large working distance between the modulator and the lens so that the lens does not block the incident light. Designing a high performance lens with a large working distance is more difficult than designing a lens of equivalent performance with no constraints on the working distance. With a transmissive spatial light modulator the working distance does not have to be long and lens design is therefore easier. In fact, as shown in Fig. 4, some transmissive spatial light modulators 31 might be useful for proximity or contact printing with no lens at all, by locating the modulator very close to the chip or wafer 34.

**[0046]** In fact, the transmissive spatial light modulator in the embodiment of Fig. 4 could be replaced by an LED array or a semiconductor laser arrays emitting light of the appropriate wavelength, each of which not only may be operated to dynamically define a desired image but also act as the light source. Thus, as modified, this embodiment would be simplified so as to not require a separate

light source.

## Claims

1. A method of deprotecting reaction sites on a substrate comprising:

providing a substrate having protected reaction sites;  
 providing a light source;  
 providing a spatial light modulator comprising an array of deformable membrane mirrors;  
 orienting said substrate, said light source, said spatial light modulator such that when said light source illuminates, intensity of illumination from said light source is modulated by said deformable membrane mirrors of said spatial light modulator and generates a predetermined light image pattern; and  
 illuminating said substrate with said predetermined light image pattern at said substrate so as to deprotect at least one of said protected reaction sites.

2. The method of claim 1, further comprising the step of projecting said predetermined light pattern onto a surface of said substrate with a lens.

3. The method of claim 2, further comprising the step of transmitting said predetermined light pattern from said lens through a micro-lens array.

4. The method of claim 2, further comprising the step of transmitting said predetermined light pattern through an array of non-imaging light concentrators.

5. The method of claim 3, further comprising the step of moving said substrate with a translation stage.

6. The method of claim 1, further comprising the step of generating a computer file that specifies, for each photolithography step, which portions of said spatial light modulator will operatively illuminate which portions of said protected reaction sites.

7. The method of claim 6, further comprising the step of programming said spatial light modulator to a desired configuration with information contained in said computer file.

8. An optical lithography system for deprotecting reaction sites on substrate, consisting essentially of:

a light source;  
 a substrate mount for mounting a substrate having protected reaction sites; and  
 a spatial light modulator for dynamically defining

a light pattern using unpatterned light from said light source without using a photomask, said spatial light modulator comprising an array of deformable membrane mirrors.

9. The optical lithography system of claim 8, wherein said spatial light modulator for dynamically defining a light pattern includes a spatial light modulator module modulating the light direction or light intensity to generate a predetermined light pattern.

10. The optical lithography systems of claim 8 or 9, wherein said light source includes an arc lamp.

11. The optical lithography system of claim 8, wherein said micro-mirror array includes a plurality of micro-mirrors, each of said micro-mirrors selectively illuminating a single feature on a substrate using specular reflection of light directed toward said substrate, and selectively not illuminating a single feature by specular reflection of light directed away from said substrate.

## Patentansprüche

1. Verfahren zum Entfernen der Schutzgruppen an Reaktionsstellen auf einem Substrat, bei dem:

ein Substrat mit geschützten Reaktionsstellen bereitgestellt wird,  
 eine Lichtquelle bereitgestellt wird,  
 ein räumlicher Lichtmodulator mit einem Feld von verformbaren Membranspiegeln bereitgestellt wird,  
 das Substrat, die Lichtquelle und der räumliche Lichtmodulator so zueinander ausgerichtet werden, dass, wenn die Lichtquelle leuchtet, die Intensität der Beleuchtung von der Lichtquelle durch die verformbaren Membranspiegel des räumlichen Lichtmodulators moduliert wird und ein vorgegebenes Licht-Bildmuster erzeugt, und  
 das Substrat mit dem vorgegebenen Licht-Bildmuster an dem Substrat beleuchtet wird, um so eine Schutzgruppenentfernung an wenigstens einer der geschützten Reaktionsstellen durchzuführen.

2. Verfahren nach Anspruch 1, bei dem weiter das vorgegebene Lichtmuster mit einer Linse auf eine Oberfläche des Substrats projiziert wird.

3. Verfahren nach Anspruch 2, bei dem man weiter das vorgegebene Lichtmuster aus der Linse ein Mikrolinsenfeld passieren lässt.

4. Verfahren nach Anspruch 2, bei dem man das vor-

gegebene Lichtmuster weiter ein Feld von nicht-ab-bildenden Lichtkonzentratoren passieren lässt.

5. Verfahren nach Anspruch 3, bei dem weiter das Substrat mit einer Verschiebungsstufe bewegt wird. 5
6. Verfahren nach Anspruch 1, bei dem weiter eine Computer-Datei erzeugt wird, die für jeden Fotolithographieschritt spezifiziert, welche Bereiche des räumlichen Lichtmodulators im Betrieb welche Bereich der geschützten Reaktionsstellen beleuchten. 10
7. Verfahren nach Anspruch 6, bei dem weiter der räumliche Lichtmodulator mit in der Computer-Datei enthaltener Information auf eine gewünschte Konfiguration programmiert wird. 15
8. Optisches Litographiesystem zur Schutzgruppenentfernung von Reaktionsstellen auf einem Substrat, das im Wesentlichen aufweist:
  - eine Lichtquelle,
  - eine Substrathalterung zum Halten eines Substrats mit geschützten Reaktionsstellen, und
  - einen räumlichen Lichtmodulator zum dynamischen Definieren eines Lichtmusters unter Verwendung von nicht-gemustertem Licht aus der Lichtquelle ohne Verwendung einer Fotomaske, wobei der räumliche Lichtmodulator ein Feld von verformbaren Membranspiegeln aufweist. 20
9. Optisches Litographiesystem nach Anspruch 8, wobei der räumliche Lichtmodulator zum dynamischen Definieren eines Lichtmusters ein räumliches Lichtmodulatorelement aufweist, das die Lichtrichtung oder Lichtintensität moduliert, um ein vorgegebenes Lichtmuster zu erzeugen. 25
10. Optisches Litographiesystem nach Anspruch 8 oder 9, wobei die Lichtquelle eine Bogenlampe umfasst. 30
11. Optisches Litographiesystem nach Anspruch 8, wobei das Mikrospiegelfeld eine Mehrzahl von Mikrospiegeln umfasst, wobei jeder Mikrospiegel selektiv ein einzelnes Merkmal auf einem Substrat beleuchtet, in dem spiegelnde Reflexion von auf das Substrat gerichtetem Licht angewendet wird, und ein einzelnes Merkmal durch spiegelnde Reflexion von dem Substrat weg gerichtetem Licht beleuchtet wird. 35

## Revendications

1. Procédé de suppression de la protection de sites de réaction sur un substrat comprenant les étapes consistant à : 40
  - fournir un substrat ayant des sites de réaction

protégés ;  
 fournir une source lumineuse ;  
 fournir un modulateur de lumière spatial comprenant un alignement de miroirs à membrane déformable ;  
 orienter ledit substrat, ladite source lumineuse, ledit modulateur de lumière spatial, de sorte que lorsque ladite source lumineuse s'illumine, l'intensité de l'illumination provenant de ladite source lumineuse est modulée par lesdits miroirs à membrane déformable dudit modulateur de lumière spatial et génère un motif d'image lumineux prédéterminé ; et  
 illuminer ledit substrat au moyen dudit motif d'image lumineux prédéterminé au niveau dudit substrat, de manière à supprimer la protection d'au moins l'un desdits sites de réaction protégés.

2. Procédé selon la revendication 1, comprenant en outre l'étape consistant à projeter ledit motif lumineux prédéterminé sur une surface dudit substrat au moyen d'une lentille. 25
3. Procédé selon la revendication 2, comprenant en outre l'étape consistant à transmettre ledit motif lumineux prédéterminé depuis ladite lentille à travers un alignement de micro-lentilles. 30
4. Procédé selon la revendication 2, comprenant en outre l'étape consistant à transmettre ledit motif lumineux prédéterminé à travers un alignement de concentrateurs de lumière non formateurs d'image. 35
5. Procédé selon la revendication 3, comprenant en outre l'étape consistant à déplacer ledit substrat au moyen d'un étage de translation. 40
6. Procédé selon la revendication 1, comprenant en outre l'étape consistant à générer un fichier informatique qui spécifie, pour chaque étape de photolithographie, quelles parties dudit modulateur de lumière spatial illumineront activement telles parties desdits sites de réaction protégés. 45
7. Procédé selon la revendication 6, comprenant en outre l'étape consistant à programmer ledit modulateur de lumière spatial dans une configuration désirée au moyen de l'information contenue dans ledit fichier informatique. 50

8. Système de lithographie optique permettant de supprimer la protection des sites de réaction sur un substrat, comprenant essentiellement : 55

une source lumineuse,  
 un support de substrat pour supporter un substrat ayant des sites de réaction protégés ; et

un modulateur de lumière spatial pour définir de façon dynamique un motif lumineux en utilisant la lumière exempte de motif provenant de ladite source lumineuse sans utiliser de masque de photogravure, ledit modulateur de lumière spatial comprenant un alignement de miroirs à membrane déformable.

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9. Système de lithographie optique selon la revendication 8, dans lequel ledit modulateur de lumière spatial permettant de définir de façon dynamique un motif lumineux comprend un module de modulateur de lumière spatial modulant la direction de la lumière ou l'intensité de la lumière pour générer un motif lumineux prédéterminé.

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10. Système de lithographie optique selon les revendications 8 ou 9, dans lequel ladite source lumineuse comprend une lampe à arc.

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11. Système de lithographie optique selon la revendication 8, dans lequel ledit alignement de micro-miroirs comprend une pluralité de micro-miroirs, chacun desdits micro-miroirs illuminant sélectivement une seule caractéristique sur un substrat en utilisant la réflexion spéculaire de la lumière dirigée vers ledit substrat, et n'illuminant pas sélectivement une seule caractéristique par réflexion spéculaire de la lumière éloignée dudit substrat.

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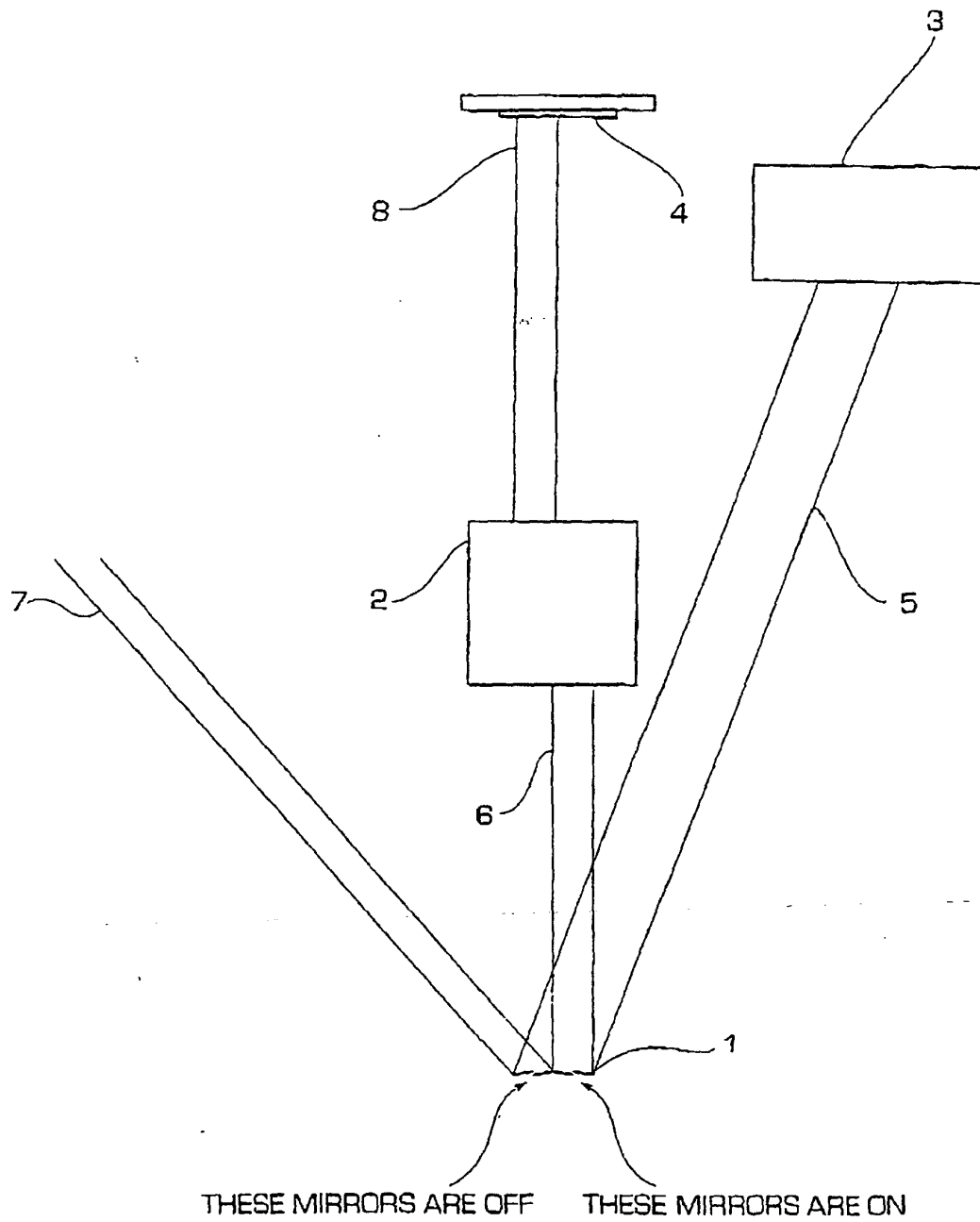


FIG. 1

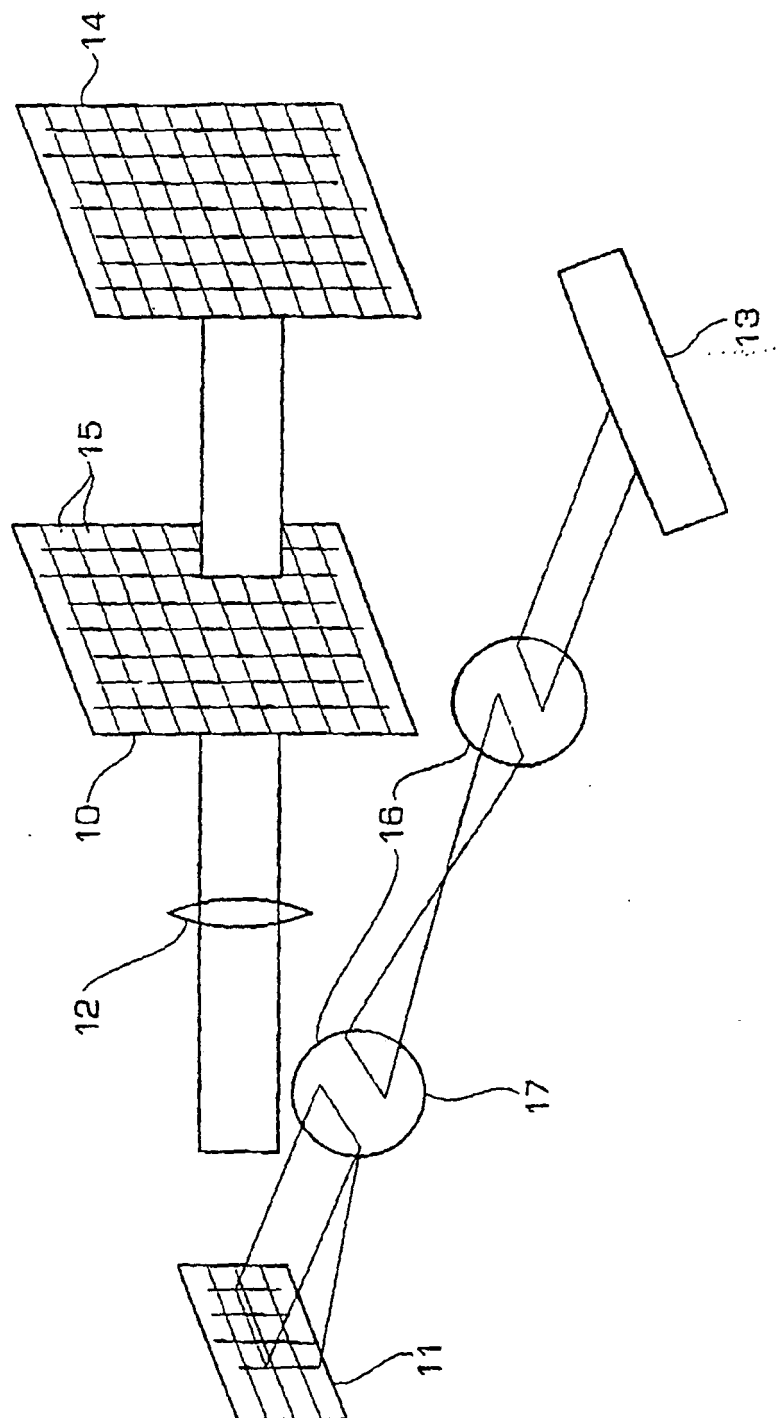


FIG. 2

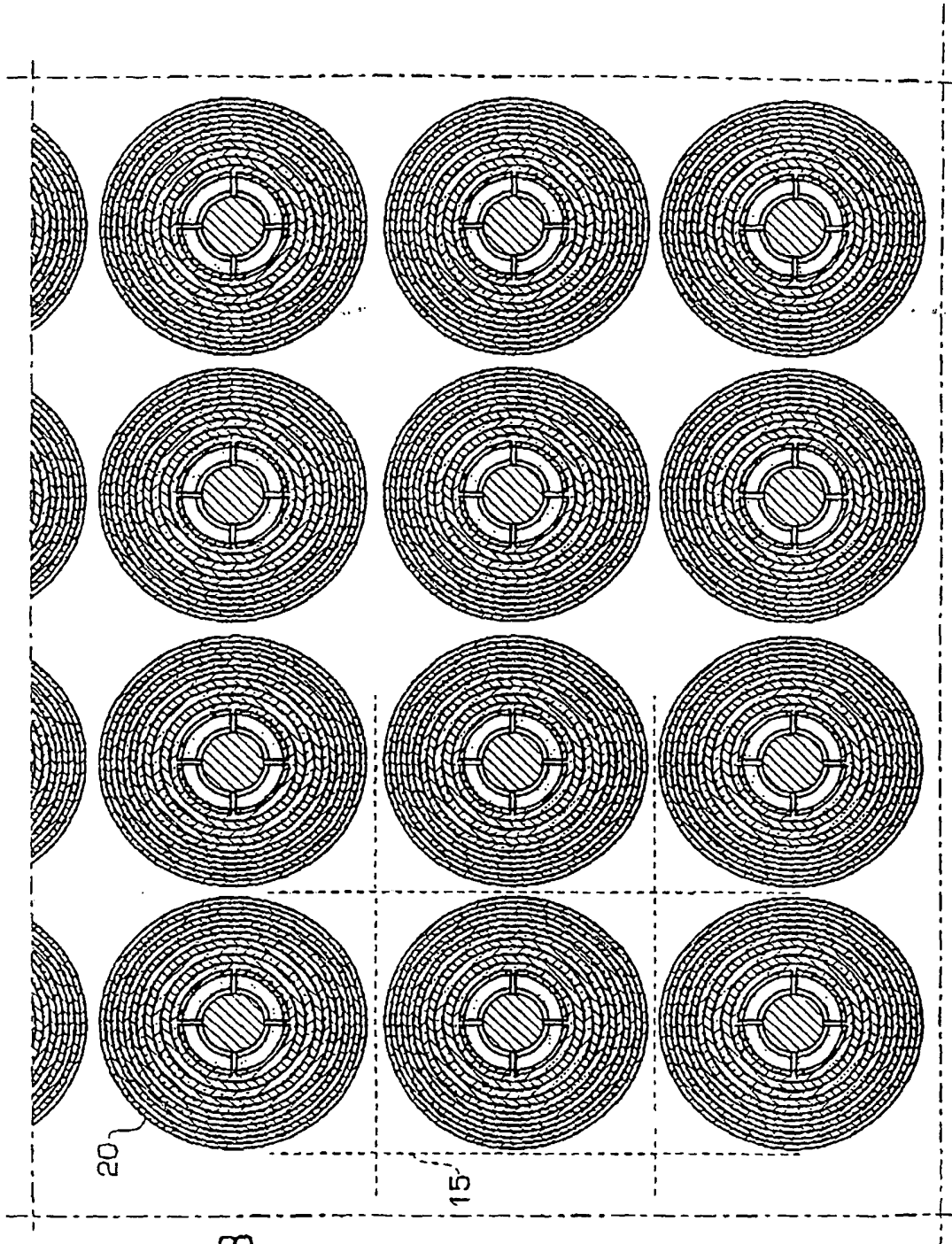


FIG. 3

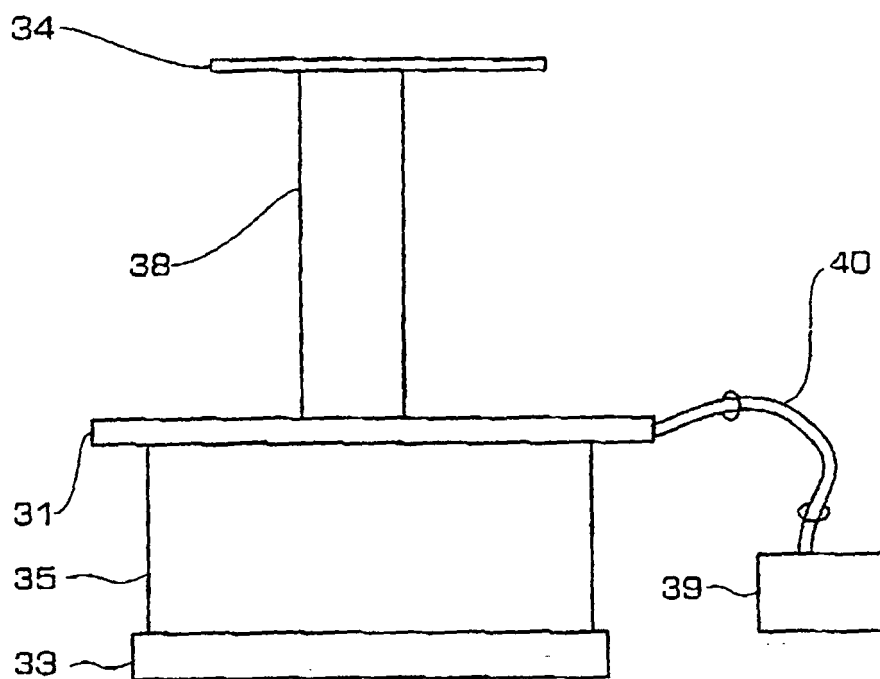


FIG. 4